

CLASSIFICATION: HEAVY CONSTANT CHAINS (cont'd)

97) MOPC173: MOUSE IGG2A
 98) CRPC101: MOUSE IGG2A
 99) IGA' CL: MOUSE IGA
 100) IGA' CL: MOUSE IGA
 101) MOPC47A: MOUSE IGA
 102) MOPC513: MOUSE IGA
 103) MOPC511: MOUSE IGA
 104) IGE' CL: MOUSE IGE
 105) IGE' CL: MOUSE IGE
 106) IGE b' CL: MOUSE IGE
 107) IGE MEMB' CL: MOUSE IGE MEMBRANE BOUND
 108) IR-731: RAT IGD
 109) RAT IGG2c' CL: RAT IGG2c
 110) RAT IGG2a' CL: RAT IGG2a
 111) RAT IGG1' CL: RAT IGG1
 112) RAT IGG2b' CL: RAT IGG2b
 113) IR2' CL: RAT IGE
 114) PM3' CL: RABBIT MU CHAIN SECRETED (ALLOTYPE A2)
 115) RABBIT IGG: RABBIT IGG
 116) PGAM6B1-12,14' CL: RABBIT IGG
 117) RABIGG: RABBIT IGG
 118) RABIGG' CL: RABBIT IGG
 119) P2A2' CL: RABBIT IGG
 120) 39-1A, 20B' CL: RABBIT IGG
 121) PA19' CL: RABBIT IGA G-SUBCLASS
 122) CT-12: COTTONTAIL RABBIT (SYLVILAGUS FLORIDANUS) IGG
 123) PIKA: PIKA (OCHOTONA RUFENSIS) IGG
 124) HA-3: HARE (LEPUS CALIFORNICUS) IGG
 125) HA-11: HARE (LEPUS CALIFORNICUS) IGG
 126) HA-1: HARE (LEPUS CALIFORNICUS) IGG
 127) HA-5: HARE (LEPUS CALIFORNICUS) IGG
 128) HA-LT: EUROPEAN HARE (LEPUF TIMIDUS) IGG
 129) SYRIAN HAMSTER IGM' CL: SYRIAN HAMSTER LSH INBRED STRAIN
 130) GP1IGG1: GUINEA PIG IGG1
 131) GP1IGG1: GUINEA PIG IGG1
 132) GP1IGG2: GUINEA PIG IGG2
 133) MOO: DOG IGM
 134) SHEEP pSHC' CL: IgG1
 135) HORSE IGG: HORSE IGG
 136) BOVINE IGG: BOVINE IGG
 137) GOAT IGG: GOAT IGG
 138) CHICKEN IGM' CL: CHICKEN IGM
 139) Ra6b' CL: ADULT Raja erinacea (LITTLE SKATE)
 140) Ra20' CL: ADULT Raja erinacea (LITTLE SKATE)
 141) Xenopus laevis IgY' CL: Xenopus laevis IGM
 142) Xenopus laevis c8(II)' CL: Xenopus laevis IGM
 143) Xenopus laevis c14(II)' CL: Xenopus laevis IGM
 144) Xenopus laevis c35' CL: Xenopus laevis IGM
 145) Xenopus laevis c40(IX)' CL: Xenopus laevis IGM
 146) Xenopus laevis J2(I)' CL: Xenopus laevis IGM
 147) Xenopus laevis J4(III)' CL: Xenopus laevis IGM
 148) Xenopus laevis J6(I)' CL: Xenopus laevis IGM
 149) Xenopus laevis J12(IV)' CL: Xenopus laevis IGM

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115) RAT IGG1'CL: BRUGGEMANN, M. (1988) GENE, 74, 473-482.

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155) Xenopus laevis cIgM'CL: SCHWAGER, J., GROSSBERGER, D. & DU PASQUIER, L. (1988) EMBO J., 7, 2409-2415.

156) Xenopus laevis J2(I)'CL: SCHWAGER, J., GROSSBERGER, D. & DU PASQUIER, L. (1988) EMBO J., 7, 2409-2415.

157) Xenopus laevis J4 (III)'CL: SCHWAGER, J., GROSSBERGER, D. & DU PASQUIER, L. (1988) EMBO J., 7, 2409-2415.

158) Xenopus laevis J6 (I)'CL: SCHWAGER, J., GROSSBERGER, D. & DU PASQUIER, L. (1988) EMBO J., 7, 2409-2415.

159) Xenopus laevis J12 (IV)'CL: SCHWAGER, J., GROSSBERGER, D. & DU PASQUIER, L. (1988) EMBO J., 7, 2409-2415.

GENERAL NOTES: HEAVY CONSTANT CHAINS

* CH1, HINGE, CH2, CH3 AND CH4 DOMAINS ARE TABULATED TO CONFORM TO THEIR CODING NUCLEOTIDE SEQUENCES RELATIVE TO INTERVENING SEQUENCES ESTABLISHED BY SAKANO, H., ROGER, J.H., HUANG, C., CHEN, J., SCHWAGER, J., MAKI, R., WALL, R. & TONEGAWA, S. (1979) NATURE, 277, 627-633. A MEMBRANE DOMAIN IS LISTED SEPARATELY TO INCLUDE PART OF THE C-TERMINAL PORTION OF MEMBRANE BOUND IMMUNOGLOBULIN. IT IS NOT NEEDED TO ALIGN THE LIGHT CHAIN WITH THE HEAVY CHAIN DOMAINS FOR HOMOLOGY. THE SEQUENTIAL NUMBERING IN THE FIRST COLUMN SHOULD BE USED: RESIDUES 108 TO 215 FOR CL; 114 TO 223 IN CH1 PLUS THE FIRST PART OF HINGE (224 TO 241); THE END OF HINGE (242 AND 243) AND THE FIRST TWO RESIDUES OF CH2 (244 AND 245); 246 THROUGH 361 OF CH2; 362 THROUGH 496 OF CH3; AND 497 THROUGH 628 OF CH4. GAPS IN THE SEQUENTIAL NUMBERING ARE USED FOR ALIGNMENT.

* DISULFIDE BONDS ARE LOCATED AT THE FOLLOWING POSITIONS IF CYS IS PRESENT:

INTRACHAIN: 142-208, 274-340, 249-312, 390-456, 524-587.

HL-INTERCHAIN: 127 OR 128, 198 OR 225, 230, 235.

HH-INTERCHAIN: 232, 233, 237, 238, 239, 240, 241, G, M, P, V, BB, EE, KK, QQ, 242, 248, 261, 314.

INTERSUBUNIT: 328, 444.

TO J-CHAIN: 495, 627.

IDENTIFICATIONS OF SOME OF THESE DISULFIDE BONDS ARE NOT ABSOLUTELY CERTAIN.

* THERE WOULD APPEAR TO BE POLYMORPHISM AMONG GAL, OU, SCO AND BOT/CO MU-CHAINS AS FOLLOWS:

GAL SER-334, VAL-358

BEST AVAILABLE COPY

SPECIFIC NOTES: HEAVY CONSTANT CHAINS

1) HUMAN IGM'CL: THE AMINO ACID SEQUENCE WAS OBTAINED BY TRANSLATING A CLONE OF HUMAN FETAL LIVER DNA.

5) BOY: FROM A CASE OF IGM HEAVY CHAIN DISEASE. THE AMINO ACID RESIDUES AT POSITIONS 451 TO 476, 519 TO 544 AND 608 TO 628 ARE IDENTICAL TO THAT OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF HUMAN DNA CONTAINING THE IGM GENE (TAKAHASHI,N., NAKAI, S. & HONJO,T. (1980) NUC.ACIDS RES., 8, 5983-5991).

7) GLI: IT HAS NO VARIABLE REGION, AND WAS FROM A CASE OF HEAVY CHAIN DISEASE.

9) WAH: THE SEQUENCES OF WAH AND NIG-65 ARE IDENTICAL FOR POSITIONS 241 PP TO 485 WHERE BOTH SEQUENCES ARE DETERMINED. FOR WAH, THERE ARE FOUR OR FIVE N-ACETYL-D-GALACTOSAMINE OLIGOSACCHARIDES ATTACHED TO SER AT POSITION 234, AND THE AZ POSITION IS 241 J, 241 K, AND 241 O AND/OR 241 P. FOR NIG-65, THERE ARE THREE N-ACETYL-D-GLUCOSAMINE OLIGOSACCHARIDES ATTACHED TO ASN AT POSITIONS 314, 414 AND 468.

10) MIG-65: THE SEQUENCES OF WAH AND NIG-65 ARE IDENTICAL FOR POSITIONS 241 PP TO 485 WHERE BOTH SEQUENCES ARE DETERMINED. FOR WAH, THERE ARE FOUR OR FIVE N-ACETYL-D-GALACTOSAMINE OLIGOSACCHARIDES ATTACHED TO SER AT POSITION 234, AND THE AZ POSITION IS 241 J, 241 K, AND 241 O AND/OR 241 P. FOR NIG-65, THERE ARE THREE N-ACETYL-D-GLUCOSAMINE OLIGOSACCHARIDES ATTACHED TO ASN AT POSITIONS 314, 414 AND 468.

11) ERI: THERE ARE THREE AMINO ACID RESIDUES BETWEEN POSITION 140 AND POSITION 142; THEY ARE ALA, VAL AND ALA. THE AUTHOR HAS RECENTLY INDICATED THAT THERE IS ONLY ONE AMINO ACID RESIDUE THERE AND IT IS ALA.

14) HUMAN IGG3'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF THREE CLONES OF HUMAN GENOMIC DNA. THERE ARE FOUR EXONS: POSITIONS 216 TO 241B, 241C TO 241O, 241R TO 241FF, AND 241GG TO 243.

15) GMN'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF HUMAN CELL LINE CDNA. FROM A CASE OF HEAVY CHAIN DISEASE.

16) HER: THE SEQUENCES OF HER, FRO AND JON ARE IDENTICAL.

17) FRO: THE SEQUENCES OF HER, FRO AND JON ARE IDENTICAL.

18) JON: THE SEQUENCES OF HER, FRO AND JON ARE IDENTICAL.

19) WIS: FROM A CASE OF HEAVY CHAIN DISEASE.

20) SPA: FROM A CASE OF HEAVY CHAIN DISEASE.

21) ZUC: FROM A CASE OF HEAVY CHAIN DISEASE.

22) ZUC': OBTAINED FROM THE SAME PATIENT AS ZUC, AND EXISTED IN A MONOMER FORM.

23) KUP: THE SEQUENCES OF KUP AND BRU ARE IDENTICAL.

24) BRU: THE SEQUENCES OF KUP AND BRU ARE IDENTICAL.

26) CHA: IT HAS NO VARIABLE REGION AND NO CH1 REGION, AND WAS FROM A CASE OF HEAVY CHAIN DISEASE. AMINO ACID RESIDUES FOUND AT POSITION 224 ARE THR AND SER.

29) NIE: IN EARLIER EDITIONS, THE AMINO ACID RESIDUES AT POSITIONS 238, 285, 300 AND 331 LISTED BY US WERE INCORRECT.

30) CRA: FROM A CASE OF HEAVY CHAIN DISEASE.

31) VAD: FROM A CASE OF HEAVY CHAIN DISEASE.

32) LKB: FROM A CASE OF HEAVY CHAIN DISEASE.

33) EST: IT HAS NO VARIABLE REGION AND NO CH1 REGION, AND WAS FROM A CASE OF HEAVY CHAIN DISEASE.

34) YOK: THE GLY AT POSITION 462 IN IGG1 CHAIN IS CONSIDERED TO BE ASSOCIATED WITH THE GM(2) ALLOTYPE (COOK,E. & STEINBERG, A.G. (1979) MOL IMMUNOL., 16, 555-558).

36) HUMAN IGG1'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A HUMAN FETAL LIVER DNA.

41) BUR: THE AMINO ACID RESIDUES AT POSITIONS 262, 263, 269, 304, 310, 333, 430, 431, 447, 461 AND 490 HAVE RECENTLY BEEN REVISED BY THE AUTHORS (PUTNAM,F.W., LIU,Y.S.V., LOW,T.L.K. (1979) J. BIOL. CHEM., 254, 2865-2874.) FROM GLN, ASP, GLX, GLN, ASN, GLN, GLY, ASP, GLU AND GLU TO GLX, ASX, GLX, GLN, ASX, GLX, GLN, ASX, GLX AND GLN RESPECTIVELY.

43) CAR: THE SEQUENCE WAS OBTAINED AFTER DIGESTION WITH PROTEASES. PROTEASES FROM S. SANGUIS AND S. PNEUMONIAE CLEAVED BETWEEN POSITIONS 241C AND 241D, AND THAT FROM H. INFLUENZAE BETWEEN POSITIONS 241G AND 241H.

47) HUMAN IGG1'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF HUMAN FETAL PSEUDO-GAMMA IGG1P. IT IS LOCATED BETWEEN IGG1A AND IGG2.

48) HUMAN IGG2'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF HUMAN GENOMIC DNA.

53) HUMAN IGG4'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF HUMAN GENOMIC DNA.

55) HUMAN IGM'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCES OF CLONES OF HUMAN GENOMIC DNA FROM FETAL LIVER (HISAMICHA,Y., MIKI,T., HISAJIMA,H. & HONJO,T. (1982) PROC.NAT.ACAD.SCI.USA, 79, 3833-3837) AND PLACENTA (MAX,E., BATTEY,J., NEY,R., KIRSCH,I.R. & HEDER,P. (1982) CELL, 29, 691-699). AND FROM A HUMAN IGM PRODUCING MYELOMA 668B1 (FLANAGAN,J.S. & RABBITS,T.H. (1982) EMBO J., 1, 655-660). ALL THREE SEQUENCES HAVE LEU AT POSITION 575 AND AN EXONIC STOP CODON 593, REPRODUCED BY AMINO ACID SEQUENCING IN IGM MYELOMA NO. MAX ET AL. AND FLANAGAN & RABBITS CARRIED OUT COMPLETE NUCLEOTIDE SEQUENCES AND FISHER ET AL. PARTIAL SEQUENCES EXCEPT FOR RESIDUE 538 WHICH IS REPORTED AS LEU BY MAX ET AL. AND AS TRP BY FLANAGAN & RABBITS; ALL AMINO ACID RESIDUES ARE IDENTICAL.

56) HUMAN IGM'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF HUMAN CDNA.

56) IGM'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF NEWBORN MOUSE DNA. THE LENGTHS OF INTERVENING SEQUENCES BETWEEN CH1 AND CH2, CH2 AND CH3, AND CH3 AND CH4 ARE 279 AND 201 BASE-PAIRS RESPECTIVELY. MATHIASSEN,S. & RABBITS,T.H. (1980) NUC.ACIDS RES., 8, 703-713. HAVE ALSO OBTAINED AMINO ACID RESIDUES 583 TO 628 FROM THE NUCLEOTIDE SEQUENCES. CALAME,K., ROGERS,E., EARLY,P., DAVIS,M., LIVANT,D., WALL,R., & HOOD,L. (1980) NATURE, 284, 452-455) BY NUCLEOTIDE SEQUENCING OF AMINO ACIDS 320 TO 343 AND 593 TO 628 FOUND ACT CODING FOR THR AT POSITION 328.

67) IGM'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE CDNA. A SINGLE AMINO ACID SUBSTITUTION AT POSITION 436 FROM SER TO ASN RESULTS IN A MUTANT PROTEIN WHICH IS DEFECTIVE IN INITIATING COMPLEMENT-DEPENDENT CYTOLYSIS. (SHULMAN,M.J., PENNELL,N., COLLINS,C. & ZOUMI,N. (1986) PROC.NAT.ACAD.SCI.USA, 83, 768-7682.)

69) MUTANT 102'CL: IT LACKS CODONS 601 TO 614, SO THAT PREDOMINANTLY MONOMERIC IgM WAS PRODUCED BY THIS CELL LINE.

70) MPC104E MEMB'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CDNA CLONE.

72) MPC76'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CDNA CLONE.

75) IGD'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE DNA.

76) IGD SEC'R'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE GENOMIC DNA.

77) IGD MEMB'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE GENOMIC DNA.

81) IGG1'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF NEWBORN MOUSE DNA. THE LENGTHS OF INTERVENING SEQUENCES BETWEEN CH1 AND HINGE, HINGE AND CH2, AND CH2 AND CH3 ARE 356, 98, AND 121 BASE-PAIRS RESPECTIVELY. HONJO,E. ET AL. HAVE DETERMINED THE ENTIRE SEQUENCE. MAX, SAKANO, T. ET AL. HAVE DETERMINED RESIDUES 316, 102 TO 268, AND 335 TO 356. THEY ARE IDENTICAL AT CORRESPONDING POSITIONS.

82) IGG1 MEMB'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE CDNA.

84) IF2'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE CDNA FROM CELL LINE IF2. THE CH1 REGION IS DELETED.

86) IGG2B(A)'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE DNA. THE LENGTHS OF INTERVENING SEQUENCES BETWEEN CH1 AND HINGE, HINGE AND CH2, AND CH2 AND CH3 ARE 316, 108, AND 112 BASE-PAIRS RESPECTIVELY.

87) IGG2B(B)'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE DNA. THE LENGTHS OF INTERVENING SEQUENCES BETWEEN CH1 AND HINGE, HINGE AND CH2, AND CH2 AND CH3 ARE 317, 109, AND 113 BASE-PAIRS RESPECTIVELY.

89) IGG2B MEMB'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE CDNA.

91) MPC11: THE AMINO ACID RESIDUES AT POSITION 363 WERE FOUND TO BE ILE AND LEU.

91) 10.1: THIS PROTEIN IS OBTAINED FROM A MUTANT OF MPC11 WITH A DELETION OF 99 NUCLEOTIDES INCLUDING THE 3' END OF THE CH1 EXON, GIVING RISE TO A DELETION OF THE ENTIRE CH1 REGION OF THE PROTEIN.

92) IGG2A(A)'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE CDNA.

93) 17/9'CL: THE FAB FRAGMENT OF 17/9 (IGG2A-KAPPA) HAS BEEN CRYSTALLIZED.

94) IGG2A(B)'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE CDNA.

95) IGG2A MEMB'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE CDNA.

96) IGM'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF BALB/C MOUSE LIVER DNA. THE SEQUENCES ARE NOT ON A SEPARATE EXON. THE FIRST THREE AMINO ACIDS 316-318 SEEM TO BE IN THE HINGE DOMAIN, AND THE REMAINING NINE AMINO ACID RESIDUES BELONG TO THE CH2 EXON. THEY HAVE BEEN PLACED IN THE HINGE DOMAIN SINCE THEY ALIGN PERFECTLY WITH OTHER IGS.

98) MPC511: CARBOHYDRATES ARE ATTACHED AT POSITIONS 154 AND 483.

104) IGM'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE DNA FROM AN ISE-PRODUCING HYBRIDOMA.

105) IGE a'CL: FROM BALB/C MOUSE LIVER DNA.

106) IGE b'CL: FROM B10.4A MOUSE LIVER DNA.

108) IGE MEMB'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF DNA FROM NEWBORN MOUSE.

111) RAT IGD'CL: THE AMINO ACID SEQUENCE WAS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF RAT CDNA.

112) IR'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF RAT CDNA.

113) IR-162'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF RAT DNA.

114) PGMMAB1-12,14'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF RABBIT CDNA.

115) RABIGG: THE AMINO ACID SEQUENCE FOR POSITIONS 266 TO 275 AND 321 TO 336 WAS ALSO CONFIRMED BY OTHERS (TEHERANI,J., CAPRA,J.D., AGGARWAL,S. & MANDY,W.J. (1978) EUR.J. IMMUNOL., 8, 690-695).

117) P2A2'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF RABBIT CDNA.

SPECIFIC NOTES: HEAVY CONS

143) SHEEP pSEC'CL: TRANSLATED

144) HORSE IGG: AS COMPARED WI

147) CHICKEN IGM'CL: THE AMINC

148) ELOPS V'CL: FROM ELOPS sa

152) XENOPUS laevis c8(XI)'CL:

153) XENOPUS laevis c14(XII)'CL:

- THE FOLLOWING WERE EQUALLY AT POSITION

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SPECIFIC NOTES: HEAVY CONSTANT CHAINS (cont'd)

143) SHEEP pSBC'CL: TRANSLATED FROM cDNA OF SHEEP LYMPHOCYTES
 144) HORSE IGG: AS COMPARED WITH HORSE IGG, THE HORSE T PROTEIN HAD VAL AT POSITION 463, GLU AT 464, AND HIS AT 474.
 147) CHICKEN IGM'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE SEQUENCE OF A CLONE OF CHICKEN cDNA.
 148) Elops VH'CL: FROM Elops saurus (LADYFISH).
 152) Xenopus laevis c8(II)'CL: ALSO KNOWN AS XIg8'CL.
 153) Xenopus laevis c14(II)'CL: ALSO KNOWN AS XIg14'CL.

+ THE FOLLOWING WERE EQUALLY AND MOST FREQUENTLY OCCURRING:

AT POSITION	RESIDUES
113D	(ILE, GLN), (ILE, GLU)
113E	(CYS, SER)
137	(THR, SER)
158	(LYS, ASN)
224	(ARG, GLU)
241B	(PRO, CYS)
281	(ALA, SER)
281	(PRO, GLU)
307	(PRO, LYS)
326	(PRO, THR)
343	(THR, ASN)
351A	(LEU, THR, ASN)
403	(THR, ASN)
452	(ASP, ASN)
496C	(PRO, ALA)
511	(ARG, GLU)
532	(ALA, GLU)
545	(PRO, LEU)
580B	(MET, ASP)
627	(LEU, VAL)
657	(GLU, GLN)
662	(ILE, VAL, ALA)
663	(ILE, SER)

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